### Amendments to the claims:

This listing of the claims will replace all prior versions, and listings of claims in the application.

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### **Listing of Claims:**

- 1. (<u>Currently Amended</u>) <u>Process A process</u> for the preparation of a statin, comprising the following steps:
  - a) Preparation of preparing a compound of the formula II

in which

S<sup>1</sup> denotes a hydrogen atom or a hydroxyl protective group,

S<sup>2</sup> and S<sup>3</sup>, independently of one another, denote hydroxyl protective groups, and

R<sup>1</sup> represents a hydrogen atom or a carboxyl protective group,

by stereoselective hydrogenation of a compound of the formula III

to give a compound of the formula II-a

$$S^3$$
 O OH O  $OR^1$  (II-a)

and optionally introduction of introducing a hydroxyl protective group; and

b) lactonization of lactonizing the compound of the formula II to give a compound of the formula I-a

- 2. (Currently Amended) Process The process according to Claim 1, comprising the further step of
- c) conversion of converting the compound of the formula I-a

into a compound of the formula I

wherein the radical

S<sup>1</sup> is as defined in Claim 1,

R denotes  $-CH_2R^2$ , -CHO,  $-CH=P(R^3)_3$ ,  $-CH_2-P^+(R^3)_3M^-$ ,

$$--CH_{2}-P-(OR^{4})_{2}$$
O
or
$$O$$

$$O$$

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R<sup>2</sup> denotes a halogen atom,  $-C \equiv N$ ,  $-CH_2NH_2$ ,  $-SO_2-R^6$  or a leaving group,

R<sup>3</sup>, R<sup>4</sup> and R<sup>5</sup> complete a Wittig radical or a Horner-Wittig radical,

denotes a hydrogen atom or a  $C_{1-3}$ -alkyl or a  $C_{5-10}$ -aryl radical, which are optionally substituted by one or more radicals which, independently of one another, are selected from halogen atoms, heterocycles which contain 0 to 10 carbon atoms and 1 to 10 heteroatoms selected from sulphur, nitrogen and oxygen

atoms, and functional groups, and

M represents an opposite ion.

3. <u>(Currently Amended) Process-The process</u> according to Claim 1-or 2, comprising the step of:

preparation of preparing a compound of the formula III

by chain extension of a compound of the formula IV

- 4. (<u>Currently Amended</u>) <u>Process-The process</u> according to <u>any of Claims 1-to 3</u>, <u>wherein the compound of the formula I being is converted into the statin by one of the following processes steps and then optionally by opening of opening the lactone ring and optionally by removal of of the protective groups:</u>
- a) reaction of reacting a compound of the formula (I)

in which the radical R represents a CHO group and the radical  $S^1$  is as defined in Claim 1, with a compound of the formula

or

in which

 $R^8$  denotes -CH=P(R<sup>3</sup>)<sub>3</sub>, -CH<sub>2</sub>-P<sup>+</sup>(R<sup>3</sup>)<sub>3</sub>M,

$$-CH_{2}-P-(OR^{4})_{2}$$
or
$$O$$

$$O$$

where R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup> and M are as defined in Claim 1,

# b) reaction of reacting a compound of the formula I

in which

the radical R denotes  $-CH=P(R^3)_3$ ,  $-CH_2-P^+(R^3)_3M^-$ ,

with a compound of the formula

$$\begin{array}{c|c}
F \\
O \\
N \\
N \\
O
\end{array}$$
or
$$\begin{array}{c|c}
R^8 \\
O \\
N \\
O
\end{array}$$
or

in which

R<sup>8</sup> denotes –CHO,

where R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup> and M are as defined in Claim 1,

## c) reaction of reacting a compound of the formula I

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in which

the radical R is a group -CH<sub>2</sub>-C $\equiv$ N,

Hydrogenation hydrogenating the compound of the formula I in which the radical R is a group -  $CH_2$ - $C\equiv N$ , to give a compound of the formula I in which the radical R is a group - $CH_2$ - $CH_2NH_2$ , and reaction of reacting the compound of the formula I in which the radical R is a group - $CH_2$ - $CH_2NH_2$  with a compound of the formula V

d) hydrogenation of hydrogenating a compound of the formula I

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#### in which

the radical R is a group - $CH_2$ - $C\equiv N$ , to give a compound of the formula I in which the radical R is a group - $CH_2$ - $CH_2NH_2$ ,

and reaction of reacting the compound of the formula I in which the radical R is a group -CH2-  $CH_2NH_2$  with a compound of the formula V

e) reaction of reacting a compound of the formula (I)

in which

the radical R is a group -CH<sub>2</sub>-CH<sub>2</sub>NH<sub>2</sub>, with a compound of the formula V

5. (Currently Amended) Process according any of Claims 1 to 4, characterized in that a compound of the formula

in which  $S^1$  is as defined in Claim 1 and St represents the radical of the statin, is converted into a compound of the formula

by catalytic hydrogenation, and optionally the protective group  $S^1$  is removed and optionally the lactone ring is opened.

6. (Currently Amended) Process-The process according to any of Claims 1-to 5, wherein the hydroxyl protective group S<sup>1</sup> being is selected from a trimethylsilyl, triisopropylsilyl, trimethylsilylethyl, tert-butyldimethylsilyl, tert-butylmethylsilyl, di-tert-butylmethylsilyl, tert-butyldiphenylsilyl, triphenylsilyl, diphenylmethylsilyl, tris(trimethylsilyl) and para-tosyl protective group.

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- 7. (Currently Amended) Process The process according to any of Claims 1-to 6, wherein the protective groups S<sup>2</sup> and S<sup>3</sup> being are bridged.
- 8. (Currently Amended) Process The process according to Claim 7, wherein the protective groups  $S^2$  and  $S^3$  together representing an isopropylidene protective group.
- 9. (Currently Amended) Process The process according to any of Claims 2-to 7, wherein the radical R representing represents a radical CH<sub>2</sub>R<sup>2</sup> and R<sup>2</sup> representing represents a leaving group, the leaving group being selected from a halogen atom, and a radical -OSO<sub>2</sub>-C<sub>1</sub>-C<sub>6</sub>-alkyl, or and -OSO<sub>2</sub>-C<sub>5</sub>-C<sub>10</sub>-aryl.
- 10. (Currently Amended) Process-The process according to any of Claims 1-to 9, wherein the radical R<sup>1</sup> denoting denotes a hydrogen atom, or a C<sub>1-3</sub>-alkyl, or a C<sub>4-10</sub>-aryl radical, each of which are may be optionally substituted by one or more radicals, which, independently of one another, are selected from halogen atoms, heterocycles which have 0 to 10 carbon atoms and 1 to 10 heteroatoms selected from sulphur, nitrogen and oxygen atoms, and functional groups.
- 11. (Currently Amended) Process The process according to any of Claims 1 to 10, wherein
- R<sup>3</sup> denoting denotes a C<sub>5</sub>- to C<sub>10</sub>-aryl radical which is optionally substituted by one or two C<sub>1</sub>-C<sub>4</sub>-alkyl radicals and/or halogen atoms, a C<sub>1</sub>-C<sub>4</sub>-alkyl radical or a C<sub>5</sub>-C<sub>10</sub>-cycloalkyl radical,
- R<sup>4</sup> denoting denotes a C<sub>1</sub>-C<sub>4</sub>-alkyl radical, and

- R<sup>5</sup> denoting denotes a C<sub>1</sub>-C<sub>6</sub>-alkyl or C<sub>5</sub>-C<sub>10</sub>-aryl radical.
- 12. <u>(Currently Amended) Process-The process</u> according to any of Claims 1-to-11, wherein the statin being is fluvastatin, rosuvastatin, cerivastatin, glenvastatin or atorvastatin.
- 13. (Currently Amended) Compound A compound of the formula I

in which

S<sup>1</sup> and R are as defined in Claim 2, with the proviso that the radical S<sup>1</sup> does not represent a tert-butyldimethylsilyl group if the radical R represents a CHO, -CH<sub>2</sub>-OTos, -CH<sub>2</sub>Cl or -CH<sub>2</sub>I group.

14. (Currently Amended) Compound according to Claim 13, in which the radical  $S^1$  represents a tert-butyldimethylsilyl group and the radical R represents a -CH<sub>2</sub>R<sup>2</sup>, -CH=P(R<sup>3</sup>)<sub>3</sub>, -CH<sub>2</sub>-P<sup>+</sup>(R<sup>3</sup>)<sub>3</sub>M<sup>-</sup>,

atom, a  $-C \equiv N$ , a  $-CH_2NH_2$  group or a radical  $-SO_2-R^6$ , and  $R^3$ ,  $R^4$ ,  $R^5$ ,  $R^6$  and M are as defined in Claim 2.

15. (Currently Amended) Process The process for the preparation of a compound of a formula (I-a)

in which the radical  $S^1$  is as defined in Claim 1, characterized in that a compound of the formula II

in which

S<sup>1</sup>, S<sup>2</sup>, S<sup>3</sup> and R<sup>1</sup> are as defined in Claim 1, is converted into the compound of the formula I-a by lactonization.